
Product Data Sheet

Product Name: Evernic Acid

Cat. No.: GC13238

Chemical Properties

Cas. No. 537-09-7

Chemical Name 2-hydroxy-4-[(2-hydroxy-4-methoxy-6-methylbenzoyl)oxy]-6-methyl-benzoic acid

SMILES CC1=CC(OC(C2=C(C)C=C(OC)C=C2O)=O)=CC(O)=C1C(O)=OFormula $C_{17}H_{16}O_7$

M.Wt 332.3

Solubility ≤ 0.5 mg/ml in ethanol; 10mg/ml in DMSO; 10mg/ml in dimethyl formamide

Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Background**

Evernic Acid is an inhibitor of two key plasmodial FAS-II enzymes PfFabZ and PfFabI

The Type II fatty acid synthase (FAS) system catalysed fatty acid biosynthesis in most bacteria. Fatty acids are aliphatic acids important for energy production and storage, cellular structure and as intermediates in the biosynthesis of hormones and other biologically important molecules. The FAS-II exists in archaea and bacteria, which has been characterized by the use of discrete, monofunctional enzymes for fatty acid synthesis. Inhibitors of this pathway (FASII) are being investigated as possible antibiotics. The bacterial FASII pathway is a promising target for the development of novel antimicrobial drugs [2]. Disruption of genes encoding mitochondrial FAS enzymes in yeast results in a respiratory-deficient phenotype and small rudimentary mitochondria [3].

Evernic acid was a secondary metabolite produced by some species of lichen. Evernic

Caution: Product has not been fully validated for medical applications. For research use only.

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acid bound to allosteric sites on the protein surface of FAS-II enzymes and produced antibacterial and antiplasmodial effects. Evernic acid inhibited PfFabZ and PfFabI with the IC₅₀ values of 10.7 and 36.1 μ M, respectively. Evernic acid showed low efficacy against the malaria parasite *P. berghei* with an IC₅₀ of 77.3 μ M [1].

References:

- [1] Lauinger I L, Vivas L, Perozzo R, et al. Potential of lichen secondary metabolites against Plasmodium liver stage parasites with FAS-II as the potential target[J]. Journal of natural products, 2013, 76(6): 1064-1070.
- [2] Marrakchi H, Zhang Y M, Rock C O. Mechanistic diversity and regulation of Type II fatty acid synthesis[J]. 2002.
- [3] Hiltunen J K, Schonauer M S, Autio K J, et al. Mitochondrial fatty acid synthesis type II: more than just fatty acids[J]. Journal of Biological Chemistry, 2009, 284(14): 9011-9015.

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